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Supplementary Information

Fourier Transform Infrared Spectroscopy was used to investigate the phase transformation in PVDF samples. The supplementary Figure 1 shows the FTIR spectra of pure PVDF and PVDF/PTFO nanocomposite samples of 5 wt% PTFO and 10 wt% PTFO. The characteristic peaks at 530, 618, 764 and 813 cm⁻¹ is a clear indication that pure PVDF as well as PVDF/PTFO nanocomposites crystallize predominantly in the α -phase. It is also observed that the addition of PTFO tends to increase the intensity of the peaks corresponding to the α -phase.



Supplementary Figure 1: FTIR analysis of pure PVDF and PVDF/PTFO nanocomposite film prepared via compression molding technique.

Supplementary Table 1: The influence of PTFO nanoparticles on PVDF matrix on the

| Parameters | PVDF/PTFO Nanocomposites | | | | | | | |
|-----------------------------------|--------------------------|----------|-----------------|----------|--|--|--|--|
| | 0 wt% | 5 wt% | 10 wt% | 25 wt% | | | | |
| E' (1kHz) | 9.7 | 9.8 | 9.7 | 12 | | | | |
| Tanδ (1kHz) | 0.0153 | 0.01182 | 0.01181 0.01098 | | | | | |
| M _s [(at 15 kOe) emu/g | - | 0.03 | 0.04 | 0.2 | | | | |
| M_r (emu/g) | - | 0.01 | 0.02 | 0.08 | | | | |
| H _c (kOe) | - | 2.83 | 3.16 | 2.68 | | | | |
| $P_{\rm m}$ (μ C/cm2) | 0.03 | 0.04 | 0.07 | 0.1 | | | | |
| $P_r (\mu C/cm2)$ | 0.01 | 0.02 | 0.023 | 0.026 | | | | |
| H _c (Kv/cm) | 3.26 | 16.12 | 17.9 | 21.22 | | | | |
| MD % | - | 0.24 | 0.26 | 0.3 | | | | |
| σ_{dc} (S/cm) | 6.07E-12 | 6.03E-12 | 6.28E-12 | 7.68E-12 | | | | |
| η | 0.96904 | 0.97069 | 0.97023 | 0.96983 | | | | |

various investigated for PVDF/PTFO nanocomposite

Supplementary Table 2: Summary of the various experimental results for pure PVDF and PVDF/PTFO nanocomposite of 25 wt% PTFO nanoparticles investigated in this work along with the relevant reported results available in the literature.

| Nanoco mpo site films | co PVDF/ PTFO ite (this work) | | Ba _{0.95} Ca _{0.05} Zr _{0.15} Ti _{0.85} O ₃ /PV DF (36) | | BaTiO ₃ : PVDF (37) | BaTi O ₃ /P VDF (38) | PVAc/ BiFeO 3 (22) | BiFeO ₃ / PVDF (19) | PVD F/BiF eO ₃ (20) | BiFe O3/ PMM A (21) | BiFeO ₃ / PVA (23) | Fe ₃ O ₄ / PVDF (39) | PVDF- GO- Fe ₃ O ₄ (40) | Ni /PVDF (41) | |
|-------------------------------------|----------------------------------|-------------|---|--------------------|--|--|---------------------------|---|---|---------------------------------|-------------------------------------|--|--|---|----------------|
| Prepera tion method | Hot m | olding | So | lution Cast | ting | Hot roll | Hot press | Soln. Cast. | Hot pres | Hot mold | Soln. Cast. | Soln. Cast. | Solv. Cast. | Solv. Cast. | Solv. Cast. |
| Filler content | 0 wt% | 25 wt% | neat PVDF | 22 vol% BCZT | 22 vol% dop- amine @ BCZT | 80 vol% BaTiO ₃ | 80% | 2wt% PureBi FeO ₃ | 50 wt% BiFeO ₃ | 22 vol% | 10 vol% | BiFeO3/ PV; 1/16 | 9.09% Fe ₃ O ₄ | 5wt% GO, 5wt% Fe ₂ O ₃ | |
| E' (1kHz) | 9.7 | 12 | ~4 | ~7 | ~9.5 | - | 70 | ~7.5 | - | 500 | ~8.5 | - | - | ~7.5 | - |
| tanð (1kHz) | 0.015 | 0.010 9 | ~0.06 | ~0.06 | ~0.07 | - | ~0.05 | ~2.2 | - | ~0.2 | ~0.05 | - | - | ~0.1 | - |
| Ms [(@ 15 kOe) emu/g | - | 0.2 | - | - | - | - | - | M _{max} ~0.4* | M _{max} ~0.02 [#] | - | ~0.6* emu/c c | ~0.5 | ~3.8 | ~2.2 | ~2.2 |
| Мг (emu/ g) ^Δ | - | 0.08 | - | - | - | - | - | - | $\sim 3.0 \times 10^{-3}$ | - | - | - | - | - | - |
| Hc (kOe)∆ | - | 2.6 | - | - | - | - | - | - | ~1.2 | - | - | - | - | - | - |
| Pm (μC/ cm ²) | 0.03 | 0.1 | - | - | - | 0.51@ 67.6 kV/ cm | ~0.8 @ 60 kV/ cm | ~4.5 at 20kV/c m | - | ~60 | - | ~0.04x1 0 ⁻³ | ~0.04 @100 kV/c m | ~0.05 at60 kV/c m | ~0.1 |
| Pr (µC/ cm ²) | 0.01 | 0.026 | - | - | - | 0.15 | ~0.2 | ~3 | - | ~20 | - | $\sim 0.02 \times 1$ 0^{-3} | ~0.02 | ~0.02 | - |
| Hc (kV/cm) | 3.26 | 21.22 | - | - | - | 19.2 | ~15k V/cm | ~15KV /cm | - | ~200 | - | 0.2kV/c m | ~50kV/ cm | ~22kV/ cm | - |
| MD % @9kOe | - | 0.3 | - | - | - | - | - | <0.10 | - | - | - | - | ~0.1 | ~0.02 | ~0.005 |
| σ _{AC} (S/cm) @ 1kHz | ~5x1 0 ⁻⁹ | ~7x10 -9 | - | - | - | - | - | - | - | ~2x 10 ⁻⁸ | ~2x10 -10 | - | - | - | - |

*samples were superparamagnetic.

samples showed no saturation.

 Δ Mr and Hc being so small that it can't be extracted from reported plots.